

## IN THE CLAIMS:

10/539054  
JC17 Rec'd PCT/PTO 15 JUN 2005

The text of all pending claims is set forth below. Cancelled and withdrawn claims are indicated with claim number and status only. The claims as listed below show added text with underlining and deleted text with ~~strikethrough~~. The status of each claim is indicated with one of (original), (currently amended). Please AMEND claims 3, 7 and 8 in accordance with the following:

1. (original): A polytrimethylene terephthalate resin comprising:

60 to 100 mole % of (a) trimethylene terephthalate recurring units, and

0 to 40 mole % of (b) at least one monomer unit selected from the group consisting of monomer units obtained from comonomers which are other than the monomers used for forming said trimethylene terephthalate recurring units and which are copolymerizable with at least one of the monomers used for forming said trimethylene terephthalate recurring units,

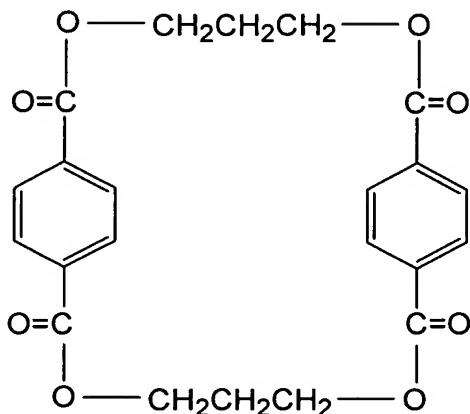
the total molar amount of (a) monomer units and (b) monomer units being 100 mole %,

said polytrimethylene terephthalate resin having the following characteristics (A) to (D):

(A) an intrinsic viscosity  $[\eta]$  of from 0.6 to 4 dl/g;

(B) a molecular weight distribution of from 2 to 2.7 in terms of the Mw/Mn ratio, wherein Mw represents the weight average molecular weight of said polytrimethylene terephthalate resin and Mn represents the number average molecular weight of said polytrimethylene terephthalate resin;

(C) a cyclic dimer content of not greater than 2 % by weight, said cyclic dimer being represented by the following formula (1):



(1); and

(D) a psychometric lightness L-value of from 70 to 100 and a psychometric chroma b\*-value of from -5 to 25.

2. (original): The polytrimethylene terephthalate resin according to claim 1, which is in the form of pellets, said pellets having a crystallinity ( $X_c$ ) of 40 % or less, wherein said crystallinity ( $X_c$ ) is defined by the following formula (2):

$$X_c (\%) = \{\rho_c \times (\rho_s - \rho_a)\} / \{\rho_s \times (\rho_c - \rho_a)\} \times 100 \quad (2)$$

wherein  $\rho_a$  is 1.300 g/cm<sup>3</sup> which is an amorphous density of trimethylene terephthalate homopolymer,  $\rho_c$  is 1.431 g/cm<sup>3</sup> which is a crystal density of trimethylene terephthalate homopolymer, and  $\rho_s$  represents a density (g/cm<sup>3</sup>) of said pellets.

3. (currently amended): A method for producing the poly-trimethylene terephthalate resin of claim 1 ~~or 2~~, which comprises:

(1) providing a crude trimethylene terephthalate resin in a molten form, said crude trimethylene terephthalate resin comprising:

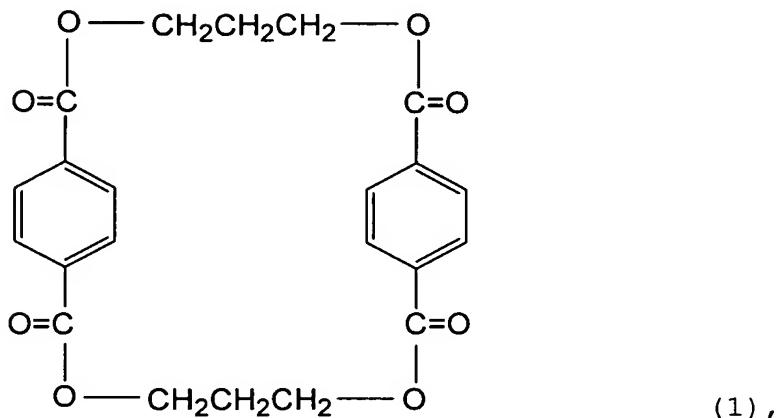
60 to 100 mole % of (a) trimethylene terephthalate recurring units, and

0 to 40 mole % of (b) at least one monomer unit selected from the group consisting of monomer units obtained from co-monomers which are other than the monomers used for forming said trimethylene terephthalate recurring units and which are copolymerizable with at least one of the monomers used for forming said trimethylene terephthalate recurring units,

the total molar amount of (a) monomer units and (b) mono-

mer units being 100 mole %,

said crude trimethylene terephthalate resin further comprising a cyclic dimer represented by the following formula (1):



said crude trimethylene terephthalate resin having an intrinsic viscosity  $[\eta]$  of from 0.2 to 4 dl/g and a cyclic dimer formation index (E) of less than 0.066, said cyclic dimer formation index (E) being defined by the following formula (3):

$$E = W/M \quad (3)$$

wherein M represents the terminal hydroxyl group content of said crude trimethylene terephthalate resin in terms of mole % based on the total molar amount of the trimethylene terephthalate unit, and W

represents the re-formation rate of the cyclic dimer in terms of an increase (as expressed by absolute percentage value) in the cyclic dimer content (% by weight), per minute, of the crude trimethylene terephthalate resin as measured at 260 °C in nitrogen gas atmosphere with respect to a sample of the crude polytrimethylene terephthalate resin in a molten form, wherein the molten sample is obtained by melting a cyclic dimer

-reduced sample of the crude polytrimethylene terephthalate resin which cyclic dimer-reduced sample has a cyclic dimer content reduced to 0.1 % by weight or less; and

(2) removing, from said crude polytrimethylene terephthalate resin in a molten form, 0.5 % by weight or more, based on the weight of said crude polytrimethylene terephthalate resin, of said cyclic dimer, by volatilization under reduced pressure.

4. (original): The method according to claim 3, wherein said crude polytrimethylene terephthalate resin has a cyclic dimer formation index (E) of less than 0.033.

5. (original): The method according to claim 3 or 4, wherein said crude trimethylene terephthalate resin in a molten form provided in said step (1) has an intrinsic viscosity  $[\eta]$  of

from 0.2 to 2 dl/g, and the removal of said cyclic dimer in said step (2) is performed by a method comprising continuously feeding said crude trimethylene terephthalate resin in a molten form to a guide-wetting fall polymerizer having a perforated plate and at least one guide provided in association with the perforated plate, wherein said crude trimethylene terephthalate resin is allowed to fall along and in contact with the surface of said at least one guide provided in the polymerizer at a temperature which is equal to or higher than the crystalline melting point of said crude trimethylene terephthalate resin and is not higher than 290 °C under reduced pressure, so that polymerization of said crude trimethylene terephthalate resin and volatilization of said cyclic dimer are effected during the fall of said crude trimethylene terephthalate resin, while continuously withdrawing the resultant trimethylene terephthalate resin from said polymerizer.

6. (original): The method according to claim 3 or 4, wherein said crude trimethylene terephthalate resin in a molten form provided in said step (1) has an intrinsic viscosity  $[\eta]$  of from 0.6 to 4 dl/g, and the removal of said cyclic dimer in said step (2) is performed by means of a thin film evaporator under conditions wherein:

(a) the pressure in said thin film evaporator is a reduced pressure of 2.6 kPa or less,

(b) a thin film of the crude polytrimethylene terephthalate resin in a molten form is formed on the inner wall of said thin film evaporator, while performing the surface renewal of said crude polytrimethylene terephthalate resin,

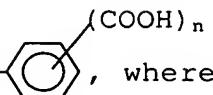
(c) said thin film of the crude polytrimethylene terephthalate resin in a molten form has a resin-gas contact area of 1 cm<sup>2</sup>/g or more, in terms of a value calculated by dividing the area of said crude polytrimethylene terephthalate resin which is in contact with the gaseous phase inside the thin film evaporator by the weight of the crude polytrimethylene terephthalate resin present in said thin film evaporator, and

(d) said crude polytrimethylene terephthalate resin occupies not more than 40 % of the inner space of said thin film evaporator.

7. (currently amended): The method according to ~~any one of~~  
~~claims 3 to 6~~ claim 3 or 4, wherein said crude trimethylene terephthalate resin is produced by a polycondensation reaction performed in the presence of a catalyst comprising at least one titanium compound and at least one phosphorus compound selected from the group consisting of phosphoric acid, a phosphoric ester, phosphorous acid, a phosphorous ester and a phosphorus compound represented by the following formula (4):



wherein  $m$  is an integer of 1 or 2, and

each  $R$  independently represents , wherein  $n$  is an integer of from 0 to 3,  
wherein said at least one titanium compound and said at least one phosphorus compound are used in respective amounts such that the phosphorus/titanium atomic ratio is in the range of from 0.01 to 10.

8. (currently amended): The method according to ~~any one of~~  
~~claims 3 to 7~~ claim 3 or 4, wherein said crude trimethylene terephthalate resin is produced by a polycondensation reaction performed in the presence of a catalyst comprising at least one tin compound having no carbon-tin bond.